Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

In the Claims:

What is claimed is:

Claims 1-15. (Cancelled).

16. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph I comprising:

- a) reacting 9-{(3aR,4R,6S,6aR)-6-[5-tert-butyl-1,3,4-oxadiazol-2-yl]-2,2-dimethyltetrahydrofuro[3,4-d][1,3]dioxol-4-yl}-N-(4-chloro-2-fluorophenyl)-9H-purin-6-amine with trifluoroacetic acid/water;
- b) neutralizing with aqueous methanolic ammonia solution at 25-50 ^oC over at least one hour; and
- c) cooling to 0-5 °C, optionally adding toluene.
- 17. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph I comprising:
- a) dissolving Compound A in N,N-dimethylformamide and water wherein the N,N-dimethylformamide:water ratio is in the range 2.5:1 to 1.5:1 and the dilution is at least 15 volumes;
- initiating crystallisation by either
 adjusting the temperature to less than 25°C; or
 adjusting the temperature to less than 30°C, and seeding with
 Polymorph I; and
- c) optionally, adding toluene.

- 18. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph II comprising:
- a) dissolving Compound A in N,N-dimethylformamide and water wherein the N,N-dimethylformamide:water ratio is in the range 2:1 to 1:2 and the dilution is at least 15 volumes;
- initiating crystallisation by
 adjusting the temperature to greater than 35°C; and optionally seeding with polymorph II; and
- c) optionally, adding toluene.
- 19. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in spheronised habit.
- 20. New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in a habit obtainable by a process of claim 16.
- 21. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in a habit obtainable by a process of claim 17.
- 22. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph II in spheronised habit.
- 23. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph II in a habit obtainable by a process of claim 18.

- 24. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 19, and a pharmaceutically acceptable carrier and/or excipient.
- 25. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 20, and a pharmaceutically acceptable carrier and/or excipient.
- 26. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 21, and a pharmaceutically acceptable carrier and/or excipient.
- 27. (New) A pharmaceutical composition comprising Polymorph II in a habit according to claim 22, and a pharmaceutically acceptable carrier and/or excipient.
- 28. (New) A pharmaceutical composition comprising Polymorph II in a habit according to claim 23, and a pharmaceutically acceptable carrier and/or excipient.
- 29. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph I of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 19.
- 30. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea

comprising administering a therapeutically effective amount of Polymorph I of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 20.

- 31. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph II of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 22.
- 32. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph II of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 23.